04/802,550

## WHAT IS CLAIMED IS:

- 1 1. An RNase A superfamily polypeptide having an N-terminus of the sequence:
- $X^1X^2SLX^3V$ , wherein  $X^1$  represents methionine or is absent,  $X^2$  represents glycine
- or is absent, and X<sup>3</sup> represents an amino acid residue, said RNase A superfamily
- 4 polypeptide being selectively toxic to a proliferating endothelial cell.
- 1 2. An RNase A superfamily polypeptide of claim 1 having SEQ. ID. No.: 2.
- 1 3. An RNase A superfamily polypeptide of claim 1 having 90% homology to SEQ.
- 2 ID. No.: 2.
- 1 4. An RNase A superfamily polypeptide of claim 1 having SEQ. ID. No.: 4.
- 1 5. An RNase A superfamily polypeptide of claim 1 having 90% homology to SEQ.
  - 2 ID. No.: 4.
- 1 6. An RNase A superfamily polypeptide of claim 1 wherein the N-terminus is
- 2 MSLHV.
- 7. An RNase A superfamily polypeptide of claim 1 wherein the N-terminus is
- 2 MGSLHV.
- 1 8. An RNase A superfamily polypeptide of claim 1 wherein the N-terminus is
- 2 attached to the EDN protein.
- 1 9. An RNase A superfamily polypeptide of claim 1 wherein the proliferating
- 2 endothelial cell is a neoplastic endothelial cell.
- 1 10. An RNase A superfamily polypeptide of claim 1 wherein the proliferating
- 2 endothelial cell is a non-neoplastic endothelial cell.
- 1 11. An RNase A superfamily polypeptide of claim 9 wherein the neoplastic
- 2 endothelial cell is a Kaposi sarcoma KS Y-1 cell.
- 1 12. An RNase A superfamily polypeptide of claim 9 wherein the neoplastic
- 2 endothelial cell is a KS Y-3 cell.

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	1	13.	An RNase A superfamily polypeptide of claim 9 wherein the neoplastic			
	2		endothelial cell is selected from the group consisting of KS 1, KS 2, KS 3, KS 4, KS 5, and KS 6 cells.			
	3					
	1 14.		A pharmaceutical composition comprising			
	2		a. a unit dosage RNase A superfamily polypeptide comprising an N-terminus			
	3		of the sequence: X <sup>1</sup> X <sup>2</sup> SLX <sup>3</sup> V, wherein X <sup>1</sup> represents methionine or is			
	4		absent, X <sup>2</sup> represents glycine or is absent, and X <sup>3</sup> represents an amino acid			
	5		residue, said RNase A superfamily polypeptide being selectively toxic to a			
	6		proliferating endothelial cell; and			
	7		b. a pharmaceutically acceptable carrier.			
the stands of th	1	15.	A method of selectively inhibiting the growth of a proliferating endothelial cell by			
	2		a. contacting said cell with an RNase A superfamily polypeptide comprising			
	3		an N-terminus of the sequence: X <sup>1</sup> X <sup>2</sup> SLX <sup>3</sup> V, wherein X <sup>1</sup> represents			
	4		methionine or is absent, X <sup>2</sup> represents glycine or is absent, and X <sup>3</sup>			
	5		represents an amino acid residue, said RNase A superfamily polypeptide			
	6		being selectively toxic to a proliferating endothelial cell; and			
	7		b. detecting the inhibition of the growth of said cell.			
	1	16.	The method of claim 15 wherein the proliferating endothelial cell is a neoplastic			
	2		cell.			
	1	17.	The method of claim 16 wherein the neoplastic cell is a Kaposi sarcoma cell.			
	1	18.	The method of claim 17 wherein the Kaposi sarcoma cell is selected from the			
	2		group consisting of KS 1, KS 2, KS 3, KS 4, KS 5, KS 6, KS Y-1, and KS Y-3			
	3		cells.			
	1	19.	A method of treating a patient with proliferating endothelial cells by			
	2		a. administering an effective amount of an RNase A superfamily polypeptide			
	3		comprising an N-terminus of the sequence: X <sup>1</sup> X <sup>2</sup> SLX <sup>3</sup> V, wherein X <sup>1</sup>			
	4		represents methionine or is absent, X <sup>2</sup> represents glycine or is absent, and			
	5		X <sup>3</sup> represents an amino acid residue, said RNase A superfamily			

polypeptide being selectively toxic to a proliferating endothelial cell; and

7 b.	detecting the a	melioration of Kaposi	sarcoma in said patient
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- 1 20. The method of claim 19 wherein the RNase A superfamily polypeptide is in an aqueous solution comprising a unit dosage and pharmaceutically acceptable excipients.
- 1 21. A method of manufacturing a pharmaceutical composition comprising the step of combining the RNase A superfamily polypeptide of claim 1 with a pharmaceutically acceptable carrier.